

EAST Search History

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	17998	esterase\$1	US-PGPUB; USPAT	AND	OFF	2006/10/02 14:00
L2	401	aquifex or pyrophilus	US-PGPUB; USPAT	AND	OFF	2006/10/02 14:00
L3	10	1 same 2	US-PGPUB; USPAT	AND	OFF	2006/10/02 14:04
L4	44	1 near5 thermostab\$	US-PGPUB; USPAT	AND	OFF	2006/10/02 14:05

* * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 14:18:14 ON 02 OCT 2006

=> fil .bec

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

0.21

0.21

FILES 'MEDLINE, SCISEARCH, LIFESCI, BIOTECHDS, BIOSIS, EMBASE, HCAPLUS, NTIS,
ESBIOBASE, BIOTECHNO, WPIDS' ENTERED AT 14:18:35 ON 02 OCT 2006
ALL COPYRIGHTS AND RESTRICTIONS APPLY. SEE HELP USAGETERMS FOR DETAILS.

11 FILES IN THE FILE LIST

=> s esterase#

FILE 'MEDLINE'

L1 24313 ESTERASE#

FILE 'SCISEARCH'

L2 15390 ESTERASE#

FILE 'LIFESCI'

L3 6257 ESTERASE#

FILE 'BIOTECHDS'

L4 2629 ESTERASE#

FILE 'BIOSIS'

L5 46005 ESTERASE#

FILE 'EMBASE'

L6 15863 ESTERASE#

FILE 'HCAPLUS'

L7 36227 ESTERASE#

FILE 'NTIS'

L8 309 ESTERASE#

FILE 'ESBIOBASE'

L9 4674 ESTERASE#

FILE 'BIOTECHNO'

L10 4672 ESTERASE#

FILE 'WPIDS'

L11 3018 ESTERASE#

TOTAL FOR ALL FILES

L12 159357 ESTERASE#

=> s aquifex or pyrophilus

FILE 'MEDLINE'

264 AQUIFEX

37 PYROPHILUS

L13 264 AQUIFEX OR PYROPHILUS

FILE 'SCISEARCH'

352 AQUIFEX

99 PYROPHILUS

L14 352 AQUIFEX OR PYROPHILUS

FILE 'LIFESCI'

175 AQUIFEX

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                28 PYROPHILUS
L15             175 AQUIFEX OR PYROPHILUS

FILE 'BIOTECHDS'
                42 AQUIFEX
                8 PYROPHILUS
L16             44 AQUIFEX OR PYROPHILUS

FILE 'BIOSIS'
                363 AQUIFEX
                58 PYROPHILUS
L17             367 AQUIFEX OR PYROPHILUS

FILE 'EMBASE'
                258 AQUIFEX
                37 PYROPHILUS
L18             258 AQUIFEX OR PYROPHILUS

FILE 'HCAPLUS'
                481 AQUIFEX
                71 PYROPHILUS
L19             481 AQUIFEX OR PYROPHILUS

FILE 'NTIS'
                1 AQUIFEX
                1 PYROPHILUS
L20             1 AQUIFEX OR PYROPHILUS

FILE 'ESBIOBASE'
                241 AQUIFEX
                34 PYROPHILUS
L21             241 AQUIFEX OR PYROPHILUS

FILE 'BIOTECHNO'
                141 AQUIFEX
                29 PYROPHILUS
L22             141 AQUIFEX OR PYROPHILUS

FILE 'WPIDS'
                39 AQUIFEX
                9 PYROPHILUS
L23             40 AQUIFEX OR PYROPHILUS

TOTAL FOR ALL FILES
L24             2364 AQUIFEX OR PYROPHILUS

=> s l12 and l24
FILE 'MEDLINE'
L25             1 L1 AND L13

FILE 'SCISEARCH'
L26             0 L2 AND L14

FILE 'LIFESCI'
L27             0 L3 AND L15

FILE 'BIOTECHDS'
L28             2 L4 AND L16

FILE 'BIOSIS'
L29             0 L5 AND L17

FILE 'EMBASE'
L30             0 L6 AND L18

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FILE 'HCAPLUS'
L31 4 L7 AND L19

FILE 'NTIS'
L32 0 L8 AND L20

FILE 'ESBIOBASE'
L33 0 L9 AND L21

FILE 'BIOTECHNO'
L34 0 L10 AND L22

FILE 'WPIDS'
L35 1 L11 AND L23

TOTAL FOR ALL FILES
L36 8 L12 AND L24

=> s l12(5a)thermostab?

FILE 'MEDLINE'
6982 THERMOSTAB?
L37 31 L1 (5A)THERMOSTAB?

FILE 'SCISEARCH'
9775 THERMOSTAB?
L38 73 L2 (5A)THERMOSTAB?

FILE 'LIFESCI'
4209 THERMOSTAB?
L39 31 L3 (5A)THERMOSTAB?

FILE 'BIOTECHDS'
7087 THERMOSTAB?
L40 62 L4 (5A)THERMOSTAB?

FILE 'BIOSIS'
10975 THERMOSTAB?
L41 60 L5 (5A)THERMOSTAB?

FILE 'EMBASE'
13453 THERMOSTAB?
L42 35 L6 (5A)THERMOSTAB?

FILE 'HCAPLUS'
20949 THERMOSTAB?
L43 108 L7 (5A)THERMOSTAB?

FILE 'NTIS'
191 THERMOSTAB?
L44 1 L8 (5A)THERMOSTAB?

FILE 'ESBIOBASE'
4084 THERMOSTAB?
L45 32 L9 (5A)THERMOSTAB?

FILE 'BIOTECHNO'
6565 THERMOSTAB?
L46 26 L10(5A)THERMOSTAB?

FILE 'WPIDS'
5360 THERMOSTAB?
L47 7 L11(5A)THERMOSTAB?

TOTAL FOR ALL FILES
L48 466 L12(5A) THERMOSTAB?

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=> s 112(10a)(gene/q or nucleic or polynucleotide#)
FILE 'MEDLINE'
    185023 NUCLEIC
    9727 POLYNUCLEOTIDE#
L49      781 L1 (10A)(GENE/Q OR NUCLEIC OR POLYNUCLEOTIDE#)

FILE 'SCISEARCH'
    37816 NUCLEIC
    4408 POLYNUCLEOTIDE#
L50      856 L2 (10A)(GENE/Q OR NUCLEIC OR POLYNUCLEOTIDE#)

FILE 'LIFESCI'
    14100 NUCLEIC
    2093 POLYNUCLEOTIDE#
L51      626 L3 (10A)(GENE/Q OR NUCLEIC OR POLYNUCLEOTIDE#)

FILE 'BIOTECHDS'
    51480 NUCLEIC
    21557 POLYNUCLEOTIDE#
L52      332 L4 (10A)(GENE/Q OR NUCLEIC OR POLYNUCLEOTIDE#)

FILE 'BIOSIS'
    54781 NUCLEIC
    7813 POLYNUCLEOTIDE#
L53      1355 L5 (10A)(GENE/Q OR NUCLEIC OR POLYNUCLEOTIDE#)

FILE 'EMBASE'
    38222 NUCLEIC
    3933 POLYNUCLEOTIDE#
L54      642 L6 (10A)(GENE/Q OR NUCLEIC OR POLYNUCLEOTIDE#)

FILE 'HCAPLUS'
    188125 NUCLEIC
    21707 POLYNUCLEOTIDE#
L55      2327 L7 (10A)(GENE/Q OR NUCLEIC OR POLYNUCLEOTIDE#)

FILE 'NTIS'
    1840 NUCLEIC
    134 POLYNUCLEOTIDE#
L56      15 L8 (10A)(GENE/Q OR NUCLEIC OR POLYNUCLEOTIDE#)

FILE 'ESBIOBASE'
    28441 NUCLEIC
    934 POLYNUCLEOTIDE#
L57      480 L9 (10A)(GENE/Q OR NUCLEIC OR POLYNUCLEOTIDE#)

FILE 'BIOTECHNO'
    19939 NUCLEIC
    1566 POLYNUCLEOTIDE#
L58      561 L10(10A)(GENE/Q OR NUCLEIC OR POLYNUCLEOTIDE#)

FILE 'WPIDS'
    63657 NUCLEIC
    26813 POLYNUCLEOTIDE#
L59      208 L11(10A)(GENE/Q OR NUCLEIC OR POLYNUCLEOTIDE#)

TOTAL FOR ALL FILES
L60      8183 L12(10A)(GENE/Q OR NUCLEIC OR POLYNUCLEOTIDE#)

=> s 148 and 160
FILE 'MEDLINE'
L61      10 L37 AND L49

FILE 'SCISEARCH'

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L62 16 L38 AND L50

FILE 'LIFESCI'

L63 8 L39 AND L51

FILE 'BIOTECHDS'

L64 19 L40 AND L52

FILE 'BIOSIS'

L65 14 L41 AND L53

FILE 'EMBASE'

L66 10 L42 AND L54

FILE 'HCAPLUS'

L67 29 L43 AND L55

FILE 'NTIS'

L68 0 L44 AND L56

FILE 'ESBIOBASE'

L69 12 L45 AND L57

FILE 'BIOTECHNO'

L70 8 L46 AND L58

FILE 'WPIDS'

L71 4 L47 AND L59

TOTAL FOR ALL FILES

L72 130 L48 AND L60

=> s l72 not 2002-2006/py

FILE 'MEDLINE'

2835793 2002-2006/PY

(20020000-20069999/PY)

L73 5 L61 NOT 2002-2006/PY

FILE 'SCISEARCH'

5217618 2002-2006/PY

(20020000-20069999/PY)

L74 5 L62 NOT 2002-2006/PY

FILE 'LIFESCI'

500908 2002-2006/PY

L75 3 L63 NOT 2002-2006/PY

FILE 'BIOTECHDS'

122128 2002-2006/PY

L76 11 L64 NOT 2002-2006/PY

FILE 'BIOSIS'

2490519 2002-2006/PY

L77 6 L65 NOT 2002-2006/PY

FILE 'EMBASE'

2456999 2002-2006/PY

L78 4 L66 NOT 2002-2006/PY

FILE 'HCAPLUS'

5379015 2002-2006/PY

L79 13 L67 NOT 2002-2006/PY

FILE 'NTIS'

71279 2002-2006/PY

L80 0 L68 NOT 2002-2006/PY

FILE 'ESBIOBASE'

1466541 2002-2006/PY

L81 6 L69 NOT 2002-2006/PY

FILE 'BIOTECHNO'

244553 2002-2006/PY

L82 6 L70 NOT 2002-2006/PY

FILE 'WPIDS'

4645299 2002-2006/PY

L83 0 L71 NOT 2002-2006/PY

TOTAL FOR ALL FILES

L84 59 L72 NOT 2002-2006/PY

=> dup rem l84

PROCESSING COMPLETED FOR L84

L85 22 DUP REM L84 (37 DUPLICATES REMOVED)

=> d tot

L85 ANSWER 1 OF 22 HCAPLUS COPYRIGHT 2006 ACS on STN

TI Stable biocatalysts for ester hydrolysis

SO U.S., 133 pp., Cont.-in-part of U.S. Ser. No. 827,810, abandoned.

CODEN: USXXAM

IN Allen, Larry; Aikens, John; Demirjian, David; Vonstein, Veronika;

Fonstein, Michael; Casadaban, Malcolm

AN 2001:279546 HCAPLUS

DN 134:307220

PATENT NO.

KIND

DATE

APPLICATION NO.

DATE

PI	US 6218167	B1	20010417	US 1998-58260	19980410
	US 6218163	B1	20010417	US 1996-694078	19960808
	US 5969121	A	19991019	US 1997-781802	19970110

L85 ANSWER 2 OF 22 LIFESCI COPYRIGHT 2006 CSA on STN DUPLICATE 1

TI Esterase isozyme polymorphism, specific and nonspecific esterase, syngenic lines development and natural occurrence of a thermostable esterase in the tropical silkworm Bombyx mori L.

SO Insect Biochemistry and Molecular Biology [Insect Biochem. Mol. Biol.], (20011100) vol. 31, no. 12, pp. 1191-1199.

ISSN: 0965-1748.

AU Chattopadhyay, G.K.; Sengupta, A.K.; Verma, A.K.; Sen, S.K.; Saratchandra, B.

AN 2002:7658 LIFESCI

L85 ANSWER 3 OF 22 BIOTECHDS COPYRIGHT 2006 THE THOMSON CORP. on STN

TI New recombinant DNA molecule comprising a sequence encoding feruloyl-esterase protein, useful for treating grasses and other plant materials used in pulp and paper industries, feed processing and food-additives;

recombinant thermostable enzyme production

AU Blum D L; Kataeva I; Li X L; Ljungdahl L G

AN 2000-07617 BIOTECHDS

PI WO 2000014243 16 Mar 2000

L85 ANSWER 4 OF 22 BIOTECHDS COPYRIGHT 2006 THE THOMSON CORP. on STN

TI New esterase, useful for manufacture of medicaments, pesticides and bioreactors;

recombinant thermostable esterase produced by

recombinant DNA technology and enzyme engineering, used to produce medicine and pesticide

AU Hourai S; Matsuki Y

AN 2000-03956 BIOTECHDS
PI EP 969094 5 Jan 2000

L85 ANSWER 5 OF 22 MEDLINE on STN DUPLICATE 2
TI Cloning, overexpression, and properties of a new thermophilic and
thermostable esterase with sequence similarity
to hormone-sensitive lipase subfamily from the archaeon *Archaeoglobus*
fulgidus.
SO Archives of biochemistry and biophysics, (2000 Jan 1) Vol. 373, No. 1, pp.
182-92.
Journal code: 0372430. ISSN: 0003-9861.
AU Manco G; Giosue E; D'Auria S; Herman P; Carrea G; Rossi M
AN 2000088609 MEDLINE

L85 ANSWER 6 OF 22 HCAPLUS COPYRIGHT 2006 ACS on STN
TI Thermally stable para-nitrobenzyl esterases
SO U.S., 112 pp.
CODEN: USXXAM
IN Arnold, Frances H.; Giver, Lorraine J.
AN 1999:561566 HCAPLUS
DN 131:181656

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5945325	A	19990831	US 1998-62890	19980420

L85 ANSWER 7 OF 22 HCAPLUS COPYRIGHT 2006 ACS on STN
TI Thermostable esterases from thermophilic bacteria and
the genes encoding them
SO PCT Int. Appl., 103 pp.
CODEN: PIXXD2
IN Allen, Larry; Aikens, John; Fonstein, Michael; Vonstein, Veronika;
Demirjian, David; Casadaban, Malcolm
AN 1998:712368 HCAPLUS
DN 129:327730

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9846770	A2	19981022	WO 1998-US7237	19980410
WO 9846770	A3	19981126		
W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
CA 2286481	AA	19981022	CA 1998-2286481	19980410
AU 9871086	A1	19981111	AU 1998-71086	19980410
EP 1005556	A2	20000607	EP 1998-918096	19980410
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
JP 2000511437	T2	20000905	JP 1998-544106	19980410

L85 ANSWER 8 OF 22 MEDLINE on STN DUPLICATE 3
TI Molecular cloning of extremely thermostable esterase
gene from hyperthermophilic archaeon *Pyrococcus furiosus* in
Escherichia coli.
SO Biotechnology and bioengineering, (1998 Mar 5) Vol. 57, No. 5, pp. 624-9.
Journal code: 7502021. ISSN: 0006-3592.
AU Ikeda M; Clark D S
AN 1999201038 MEDLINE

L85 ANSWER 9 OF 22 MEDLINE on STN DUPLICATE 4
TI Overexpression and properties of a new thermophilic and
thermostable esterase from *Bacillus acidocaldarius* with

sequence similarity to hormone-sensitive lipase subfamily.
SO The Biochemical journal, (1998 May 15) Vol. 332 (Pt 1), pp. 203-12.
Journal code: 2984726R. ISSN: 0264-6021.
AU Manco G; Adinolfi E; Pisani F M; Ottolina G; Carrea G; Rossi M
AN 1998244829 MEDLINE

L85 ANSWER 10 OF 22 BIOTECHDS COPYRIGHT 2006 THE THOMSON CORP. on STN
TI A novel heat-stable lipolytic enzyme from *Sulfolobus acidocaldarius* DSM
639 displaying similarity to polyhydroxyalkanoate-depolymerases;
esterase Est1 gene cloning and enzyme
characterization
SO FEMS Microbiol.Lett.; (1998) 167, 1, 69-73
CODEN: FMLED7 ISSN: 0378-1097
AU Arpigny J L; Jendrosseck D; Jaeger K E
AN 1999-11789 BIOTECHDS

L85 ANSWER 11 OF 22 BIOTECHDS COPYRIGHT 2006 THE THOMSON CORP. on STN
TI New nucleic acid encoding heat stable esterases from
thermophilic bacteria;
recombinant thermostable esterases for use in
pharmaceutical, agricultural or food industries, etc.
AU Robertson D E; Murphy D; Reid J; Maffia A M; Link S; Swanson R V; Warren
P V; Kosmotka A; Callen W
AN 1997-11973 BIOTECHDS
PI WO 9730160 21 Aug 1997

L85 ANSWER 12 OF 22 HCAPLUS COPYRIGHT 2006 ACS on STN
TI Cloning of cDNA for and preparation of stereoselective,
thermostable esterase of *Klebsiella oxytoca*
SO Jpn. Kokai Tokkyo Koho, 14 pp.
CODEN: JKXXAF
IN Nomoto, Shiki; Kuramura, Akiko; Utsura, Kensaku
AN 1997:705999 HCAPLUS
DN 128:20050

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 09275982	A2	19971028	JP 1996-91571	19960412

L85 ANSWER 13 OF 22 MEDLINE on STN DUPLICATE 5
TI Isolation, analysis, and expression of two genes from
Thermoanaerobacterium sp. strain JW/SL YS485: a beta-xylosidase and a
novel acetyl xylan esterase with cephalosporin C deacetylase activity.
SO Journal of bacteriology, (1997 Sep) Vol. 179, No. 17, pp. 5436-41.
Journal code: 2985120R. ISSN: 0021-9193.
AU Lorenz W W; Wiegel J
AN 97431493 MEDLINE

L85 ANSWER 14 OF 22 HCAPLUS COPYRIGHT 2006 ACS on STN
TI Purification and some properties of a thermostable acid
esterase from *Acidiphilium* sp. AIU 409
SO Journal of General and Applied Microbiology (1997), 43(3), 151-156
CODEN: JGAMA9; ISSN: 0022-1260
AU Isobe, Kimiyasu; Wakao, Norio
AN 1997:628371 HCAPLUS
DN 127:216848

L85 ANSWER 15 OF 22 BIOTECHDS COPYRIGHT 2006 THE THOMSON CORP. on STN
TI A thermostable esterase;
mutant enzyme produced by enzyme engineering
AN 1995-14811 BIOTECHDS
PI JP 07213280 15 Aug 1995

L85 ANSWER 16 OF 22 MEDLINE on STN DUPLICATE 6
TI Nucleotide sequence of the gene for a
thermostable esterase from *Pseudomonas putida* MR-2068.

SO Bioscience, biotechnology, and biochemistry, (1995 Jul) Vol. 59, No. 7,
pp. 1204-7.
Journal code: 9205717. ISSN: 0916-8451.

AU Ozaki E; Sakimae A; Numazawa R
AN 95399993 MEDLINE

L85 ANSWER 17 OF 22 BIOTECHDS COPYRIGHT 2006 THE THOMSON CORP. on STN
TI A DNA fragment having a DNA sequence coding esterase;
recombinant thermostable esterase production
AN 1994-08916 BIOTECHDS
PI JP 06105693 19 Apr 1994

L85 ANSWER 18 OF 22 HCAPLUS COPYRIGHT 2006 ACS on STN
TI Novel esterase of Bacillus stearothermophilus
SO Jpn. Kokai Tokkyo Koho, 6 pp.
CODEN: JKXXAF
IN Yamane, Tsuneo; Ueda, Shunsaku; Amagi, Jusuke; Kugimya, Wataru; Takagi,
Hiroaki
AN 1994:528768 HCAPLUS
DN 121:128768

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 06165675	A2	19940614	JP 1992-5037	19920114

L85 ANSWER 19 OF 22 BIOTECHDS COPYRIGHT 2006 THE THOMSON CORP. on STN
TI DNA sequence encoding an esterase;
used for stereospecific hydrolysis of carboxylic acid ester to give
optically active acid; Pseudomonas putida gene cloning in Escherichia
coli
AN 1993-01116 BIOTECHDS
PI EP 513806 19 Dec 1992

L85 ANSWER 20 OF 22 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on
STN
TI PURIFICATION AND PROPERTIES OF EXTRACELLULAR LIPASE FROM
PSEUDOMONAS-AERUGINOSA EF2.
SO Journal of General Microbiology, (1991) Vol. 137, No. 9, pp. 2223-2230.
CODEN: JGMIAN. ISSN: 0022-1287.
AU GILBERT E J [Reprint author]; CORNISH A; JONES C W
AN 1992:32019 BIOSIS

L85 ANSWER 21 OF 22 BIOTECHDS COPYRIGHT 2006 THE THOMSON CORP. on STN
TI Overproduction of an acetyl-xylan-esterase from the extreme thermophile
'Caldocellum saccharolyticum' in Escherichia coli;
xynC gene cloning; plasmid pNZ1600 and plasmid pNZ1447 construction
SO Appl.Microbiol.Biotechnol.; (1990) 34, 2, 214-19
CODEN: EJABDD
AU Luthi E; Jasmat N B; *Bergquist P L
AN 1991-04703 BIOTECHDS

L85 ANSWER 22 OF 22 HCAPLUS COPYRIGHT 2006 ACS on STN
TI Studies of esterase-6 in Drosophila melanogaster. II. The genetics and
frequency distributions of naturally occurring variants studied by
electrophoretic and heat stability criteria
SO Genetics (1979), 93(2), 461-78
CODEN: GENTAE; ISSN: 0016-6731
AU Cochrane, Bruce J.; Richmond, Rollin C.
AN 1980:125432 HCAPLUS
DN 92:125432

=> d ab 4,5,7,14-19

L85 ANSWER 4 OF 22 BIOTECHDS COPYRIGHT 2006 THE THOMSON CORP. on STN
AB An esterase consisting of at least part of the given 370 amino acid

protein sequence with thermostable esterase activity, is claimed. It may be modified by one or more of the following amino acid substitutions: amino acid 435 substituted by isoleucine; amino acid 240 substituted with alanine; or amino acid 43 substituted with serine. Also claimed is a gene encoding the esterase, a plasmid containing that gene, a host cell transformed by the plasmid, and a means of producing the esterase. These can be used to produce medicines, pesticides and intermediates used to produce them. The enzyme can also be used in ester hydrolysis, synthesis and interchange. The esterase has a high thermostability and shortens the reaction time and promotes the reaction efficiency of ester synthesis and interchange. The host cell may be a yeast, mammal or prokaryotic cell, especially a *Chromobacterium*. The nucleic acid encoding the esterase has a given 1,110 bp DNA sequence. The plasmid used to express that gene is specifically plasmid pCC43S. (30pp)

L85 ANSWER 5 OF 22 MEDLINE on STN DUPLICATE 2

AB A new esterase gene from the hyperthermophilic archaeon *Archaeoglobus fulgidus*, reported to show homology with the mammalian hormone-sensitive lipase (HSL)-like group of the esterase/lipase family, was cloned by means of the polymerase chain reaction from the *A. fulgidus* genome. In order to compare the biochemical properties of this putative hyperthermophilic enzyme with those of the homologous, thermophilic member of HSL group, namely *Alicyclobacillus* (formerly *Bacillus*) *acidocaldarius* esterase 2 (EST2), an overexpression system in *Escherichia coli* was established. The recombinant protein, expressed in soluble and active form at 20 mg/liter of *E. coli* culture, was purified to homogeneity and characterized. The enzyme, a 35.5-kDa monomeric protein, was demonstrated to be a B"-type carboxylesterase (EC 3.1.1.1) on the basis of substrate specificity and the action of inhibitors. Among the p-nitrophenyl (PNP) esters tested the best substrate was PNP-hexanoate with $K(m)$ and $k(cat)$ values of $11 \pm 3 \mu M$ (mean \pm SD, $n = 3$) and $1014 \pm 38 s^{-1}$ (mean \pm SD, $n = 3$), respectively, at 70 degrees C and pH 7.1. Inactivation by diethylpyrocarbonate, phenylmethylsulfonylfluoride, diisopropylfosfofluoridate (DFP), and physostigmine, as well as labeling with $[(3)H]DFP$, supported our previous suggestion of a catalytic triad made up of Ser(160)-His(285)-Asp(255). The sequence identity with the thermostable *A. acidocaldarius* EST2 was 42.5%. The enzyme proved to be much more stable than its *Alicyclobacillus* counterpart. The conformational dynamics of the two proteins were investigated by frequency-domain fluorometry and anisotropy decay and the activity/stability/temperature relationship was discussed. Copyright 2000 Academic Press.

L85 ANSWER 7 OF 22 HCAPLUS COPYRIGHT 2006 ACS on STN

AB Novel thermostable esterases for industrial use are identified in thermophilic bacteria. Methods and kits for identifying thermostable esterases and for quickly determining their patterns of substrate use are described. The enzymes are characterized and genes encoding them are cloned and expressed. Two esterases (E100 and E101) were identified in *Thermus* sp. T351. The two enzymes had different substrate preferences but were both inhibited by PMSF. A total of 20 thermostable esterases were identified in a number of incompletely characterized thermophilic bacteria. Most of the enzymes had a temperature optimum of 45° and were active at near-neutral pH's. Genes were cloned by expression from a Sau3A partial digest library in λ ZAP by overlaying phage plates with an agar containing a chromogenic esterase substrate.

L85 ANSWER 14 OF 22 HCAPLUS COPYRIGHT 2006 ACS on STN

AB Extracellular and cell-bound esterases produced by *Acidiphilium* sp. AIU 409 were homogeneously purified from culture broth and cells, resp., and some properties were investigated. Both esterases more rapidly hydrolyzed p-nitrophenyl acyl esters containing long-chain fatty acids from C8:0 to C18:0

than those containing short-chain fatty acids from C2:0 to C6:0. The Km values for p-nitrophenyl long-chain fatty acid esters from C8:0 to C18:0 were approx. 1.3-1.5 mM. The enzymes were stable at 50° for 2 days between pH 3.0 and 6.5, and the optimum pH and temperature were 5.0 and 70°, resp. Enzyme activity was inhibited by phenylmethylsulfonyl fluoride and SDS. The mol. weight of both enzymes was estimated to be .apprx.64

kDa by SDS-PAGE. The N-terminal sequence was the same in both enzymes. The results suggested that extracellular esterase might be composed of the same components as cell-bound esterase.

L85 ANSWER 15 OF 22 BIOTECHDS COPYRIGHT 2006 THE THOMSON CORP. on STN
AB A thermostable esterase is claimed containing a site-specific mutation at amino acid 160 and/or 189 of a specified protein sequence; a partial amino acid sequence represents the thermostable esterase activity. Also claimed are: a gene encoding the esterase; a plasmid encoding the gene; a microorganism transformed with the plasmid; and the preparation of a mutant esterase in which the transformed microorganism is cultured for the production of the esterase. The recombinant, thermostable esterase is produced efficiently, and can be used in organic synthesis. In an example, *Chromobacterium* sp. SC-YM-1 was cultured in a medium containing 1% glucose, 1% yeast extract, 0.1% K₂HPO₄ and 0.02% MgSO₄ at 30 deg. When the OD₆₆₀ reached 3.4, 2 U/ml of benzylpenicillin was added to the culture, which was continued until the OD₆₆₀ reached 10. The microorganism was recovered by centrifugation, mixed with Tris buffer, sucrose and lysozyme (EC-3.2.1.17), and incubated at 37 deg for 30 min. SDS and protease-K were added and the mixture was incubated for a further 3 hr. Microorganism DNA was purified and used in the construction of a DNA library, which was screened and subcloned. (16pp)

L85 ANSWER 16 OF 22 MEDLINE on STN DUPLICATE 6
AB The esterase gene (est) of *Pseudomonas putida* MR-2068 was cloned into *Escherichia coli* JM109. An 8-kb inserted DNA directed synthesis of an esterase in *E. coli*. The esterase gene was in a 1.1-kb PstI-ClaI fragment within the insert DNA. The complete nucleotides of the DNA fragment containing the esterase gene were sequenced and found to include a single open reading frame of 828 bp coding for a protein of 276 amino acid residues. The open reading frame was confirmed by N-terminal amino acid sequence analysis of the purified esterase. A potential Shine-Dalgarno sequence is followed by the open reading frame. The esterase activity of the recombinant *E. coli* was more than 200 times higher than that of parental strain, *P. putida* MR-2068.

L85 ANSWER 17 OF 22 BIOTECHDS COPYRIGHT 2006 THE THOMSON CORP. on STN
AB A DNA fragment of a specified sequence coding an esterase capable of catalyzing the asymmetric hydrolysis of a carboxylic acid ester of formula (I), where R₁ = alkyl, aralkyl or aryl, R₂ and R₃ = alkyl, n = 1 or 2, is claimed. The DNA fragment is useful for the large-scale preparation of a thermostable esterase. In an example, *Pseudomonas putida* MR-2068 chromosomal DNA was digested and ligated with plasmid pUC19. *Escherichia coli* JM109 was reacted with the ligand and strains having recombinant plasmids containing the inserted structural esterase gene were selected. The transformant was cultured and the recombinant plasmid was designated plasmid pPE101. The esterase structural gene DNA sequence was determined. The thermostabilities of *Pseudomonas fluorescens* IFO 3018, *Pseudomonas putida* MR-2068 and *Escherichia coli* JM109 harboring plasmid pPE116 were examined. The remaining esterase activities after treatment at 70 deg for 3 hr were 30, 100 and 100%, respectively. (13pp)

L85 ANSWER 18 OF 22 HCAPLUS COPYRIGHT 2006 ACS on STN

AB A novel thermostable esterase is produced by *B. brevis* transformed with the esterase-encoding gene of *B. stearothermophilus*. The enzyme exhibits a pH optimum 7.5, a temperature optimum 55°, and a mol. weight of 29,000 on SDS-PAGE. The esterase is highly specific to the esters containing lower fatty acids.

L85 ANSWER 19 OF 22 BIOTECHDS COPYRIGHT 2006 THE THOMSON CORP. on STN
 AB A DNA fragment contains a DNA sequence that encodes an esterase. The esterase is capable of asymmetric hydrolysis of carboxylic acid esters of formula $R_1-COS-(CH_2)_n-CHR_2-COOR_3$ to give optically active carboxylic acids of formula $R_1-COS-(CH_2)_n-CHR_2-COOH$. In these formulae, R_1 = alkyl, aralkyl or aryl, R_2 and R_3 = alkyl, and n = 1 or 2. Also claimed are: the esterase; a recombinant plasmid including all or part of the claimed DNA fragment; a bacterial transformant carrying the plasmid; and a method of producing the optically active carboxylic acid and its enantiomer ester by reacting a racemic mixture of carboxylic acid esters with cells of the recombinant bacterium or its cellular products. To obtain the recombinant esterase, chromosomal DNA was prepared from *Pseudomonas putida* FERM BP-3846 and cloned into plasmid pUC19, and used to transform *Escherichia coli* JM105 to obtain transformants containing DNA coding for esterase gene expression. The esterase is stable up to 70 deg, the optimum temperature being 60-70 deg. Transformants containing the esterase gene have higher activity than the donor because they contain a high copy number plasmid. (27pp)

=> log y

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

124.24

124.45

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE

TOTAL

ENTRY

SESSION

CA SUBSCRIBER PRICE

-2.25

-2.25

STN INTERNATIONAL LOGOFF AT 15:31:55 ON 02 OCT 2006